

```

is in DialUnits
? b 410
    01jul10 12:29:18 User208760 Session D3196.1
        $0.57    0.152 DialUnits File1
$0.57 Estimated cost File1
$0.02 TELNET
$0.59 Estimated cost this search
$0.59 Estimated total session cost    0.152 DialUnits

```

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File 410:The Chronolog 1991-2010/ Mar
(c) 2010 Dialog. All rights reserved.

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Set Items Description
---
? set hi ;set hi
HIGHLIGHT set on as ''
HIGHLIGHT set on as ''
? begin 5,73,155,399
    01jul10 12:29:25 User208760 Session D3196.2
        $0.00    0.115 DialUnits File410
$0.00 Estimated cost File410
$0.03 TELNET
$0.03 Estimated cost this search
$0.62 Estimated total session cost    0.267 DialUnits

```

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SYSTEM:OS - DIALOG OneSearch
File 5:Biosis Previews(R) 1926-2010/Jun W4
(c) 2010 The Thomson Corporation
File 73:EMBASE 1974-2010/Jul 01
(c) 2010 Elsevier B.V.
*File 73: The archive of Medline derived records was added to Embase.
File 155:MEDLINE(R) 1950-2010/Jun 29
(c) format only 2010 Dialog
*File 155: Medline has been reloaded. Please see HELP NEWS154
for information.
File 399:CA SEARCH(R) 1967-2010/UD=15301
(c) 2010 American Chemical Society
*File 399: Use is subject to the terms of your user/customer agreement.
IPCR/8 classification codes now searchable as IC=. See HELP NEWSIPCR.

```

```

Set Items Description
---
? e au=tedesco francesco ?

Ref Items Index-term
E1 4 AU=TEDESCO F.S.
E2 101 AU=TEDESCO FRANCESCO
E3 0 *AU=TEDESCO FRANCESCO ?
E4 6 AU=TEDESCO FRANCESCO SAVERIO
E5 32 AU=TEDESCO G
E6 28 AU=TEDESCO G.
E7 2 AU=TEDESCO G.D.
E8 1 AU=TEDESCO GIANNI
E9 1 AU=TEDESCO GIOACCHINO
E10 31 AU=TEDESCO GIOVANNA
E11 2 AU=TEDESCO GISELLE DAHAREM
E12 4 AU=TEDESCO GISELLE DARAHAM

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Enter P or PAGE for more
? s e1-e4
    4 AU=TEDESCO F.S.

```

101 AU=TEDESCO FRANCESCO
 0 AU=TEDESCO FRANCESCO ?
 6 AU=TEDESCO FRANCESCO SAVERIO
 S1 111 E1-E4
 ? e au=marzari roberto ?

| Ref | Items | Index-term |
|-----|-------|----------------------------|
| E1 | 63 | AU=MARZARI R. |
| E2 | 72 | AU=MARZARI ROBERTO |
| E3 | 0 | *AU=MARZARI ROBERTO ? |
| E4 | 3 | AU=MARZARI VICTOR |
| E5 | 1 | AU=MARZARI-CHIESA A |
| E6 | 1 | AU=MARZARI-CHIESA A. |
| E7 | 95 | AU=MARZARI-CHIESA, A. |
| E8 | 3 | AU=MARZARI-CHIESA, ALBERTA |
| E9 | 3 | AU=MARZARI, A. |
| E10 | 1 | AU=MARZARI, C. |
| E11 | 1 | AU=MARZARI, CHIARA |
| E12 | 1 | AU=MARZARI, CHRISTIAN M. |

Enter P or PAGE for more

? s e1-e2
 63 AU=MARZARI R.
 72 AU=MARZARI ROBERTO
 S2 135 E1-E2
 ? s (s1 or s2) and (antibod? or immunoglobulin? or hybridoma?) (20n) (c5a or ts(w)al2?)
 111 S1
 135 S2
 2724863 ANTIBOD?
 1064219 IMMUNOGLOBULIN?
 61953 HYBRIDOMA?
 14523 C5A
 50236 TS
 5903 Al2?
 4 TS(W)Al2?
 1785 ((ANTIBOD? OR IMMUNOGLOBULIN?) OR HYBRIDOMA?) (20N) (C5A OR TS(W)Al2?)
 S3 5 (S1 OR S2) AND (ANTIBOD? OR IMMUNOGLOBULIN? OR HYBRIDOMA?) (20N) (C5A OR TS(W)Al2?)
 ? rd s3
 S4 2 RD S3 (unique items)
 ? t s4/3/all

4/3/1 (Item 1 from file: 5)
 DIALOG(R)File 5:Biosis Previews(R)
 (c) 2010 The Thomson Corporation. All rts. reserv.

16983076 BIOSIS NO.: 200200576587

The cleavage site of C5 from man and animals as a common target for neutralizing human monoclonal antibodies: In vitro and in vivo studies
 AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti Fabio; Gennaro Renato; Marks James D; Bradbury Andrew; Tedesco Francesco (Reprint)

AUTHOR ADDRESS: Dipartimento di Fisiologia e Patologia, Universita di Trieste, Via Fleming 22, I-34127, Trieste, Italy**Italy

JOURNAL: European Journal of Immunology 32 (10): p2773-2782 October, 2002
 2002

MEDIUM: print

ISSN: 0014-2980

DOCUMENT TYPE: Article

RECORD TYPE: Abstract
LANGUAGE: English

4/3/2 (Item 1 from file: 73)
DIALOG(R)File 73:EMBASE
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0081757282 EMBASE/Medline No: 2007191236
Selective therapeutic control of C5a and the terminal complement complex
by anti-C5 single-chain Fv in an experimental model of antigen-induced
arthritis in rats
Fischetti F.; Durigutto P.; Macor P.; ***Marzari R.*** ; Carretta R.;
Tedesco F.
University of Trieste, Trieste, Italy
AUTHOR EMAIL: tedesco@units.it
CORRESP. AUTHOR/AFFIL: Tedesco F.: Department of Physiology and
Pathology, University of Trieste, via Valerio 28, 34127 Trieste, Italy
CORRESP. AUTHOR EMAIL: tedesco@units.it

Arthritis and Rheumatism (Arthritis Rheum.) (United States) April 1,
2007, 56/4 (1187-1197)
CODEN: ARHEA ISSN: 0004-3591
DOI: 10.1002/art.22492
DOCUMENT TYPE: Journal; Article RECORD TYPE: Abstract
LANGUAGE: English SUMMARY LANGUAGE: English
NUMBER OF REFERENCES: 56
? t s4/7/all

4/7/1 (Item 1 from file: 5)
DIALOG(R)File 5:BIOSIS Previews(R)
(c) 2010 The Thomson Corporation. All rts. reserv.

16983076 BIOSIS NO.: 200200576587
The cleavage site of C5 from man and animals as a common target for
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AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti
Fabio; Gennaro Renato; Marks James D; Bradbury Andrew; Tedesco
Francesco (Reprint)
AUTHOR ADDRESS: Dipartimento di Fisiologia e Patologia, Università di
Trieste, Via Fleming 22, I-34127, Trieste, Italy**Italy
JOURNAL: European Journal of Immunology 32 (10): p2773-2782 October, 2002
2002
MEDIUM: print
ISSN: 0014-2980
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English

ABSTRACT: The isolation of an anti-C5 single-chain fragment variable (scFv)
antibody, TS-A12/22, from a human phage display library, is
described. This ***antibody*** inhibits the activation of C5 and the
assembly of the terminal complement complex implicated in cell and tissue
damage. Using ***antibody*** -sensitized sheep erythrocytes and rabbit red
cells as target cells in hemolytic assays, we found that TS-
A12/22 inhibited the activation of C5 by the convertases of both
classical and alternative pathways. Western blot analysis and competition
experiments with synthetic peptides showed that TS-A12/22 reacted with
the a chain of C5 and recognized the cleavage site of this complement
component by the C5 convertase. As a result, the ***antibody*** prevented
splitting of C5 and inhibited the generation of C5a and of the

terminal complement complex. The identification of the ***TS*** - ***A12*** /22 recognition site as a conserved sequence in man, mouse, rat and rabbit enabled the demonstration of in vitro inhibition of complement activity in these species. The scFv TS-A12/22 was tested in a rat model of antigen-induced arthritis and proved to be effective in preventing influx of polymorphonuclear cells into the knee joint and C9 deposition on synovial tissue.

4/7/2 (Item 1 from file: 73)
DIALOG(R)File 73:EMBASE
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Selective therapeutic control of C5a and the terminal complement complex by anti-C5 single-chain Fv in an experimental model of antigen-induced arthritis in rats

Fischetti F.; Durigutto P.; Macor P.; ***Marzari R.*** ; Carretta R.; Tedesco F.

University of Trieste, Trieste, Italy

AUTHOR EMAIL: tedesco@units.it

CORRESP. AUTHOR/AFFIL: Tedesco F.: Department of Physiology and

Pathology, University of Trieste, via Valerio 28, 34127 Trieste, Italy

CORRESP. AUTHOR EMAIL: tedesco@units.it

Arthritis and Rheumatism (Arthritis Rheum.) (United States) April 1, 2007, 56/4 (1187-1197)

CODEN: ARHEA ISSN: 0004-3591

DOI: 10.1002/art.22492

DOCUMENT TYPE: Journal; Article RECORD TYPE: Abstract

LANGUAGE: English SUMMARY LANGUAGE: English

NUMBER OF REFERENCES: 56

Objective. To determine the role of the terminal complement complex (TCC) in the development of experimental antigen-induced arthritis (AIA) and the therapeutic effects of human anti-C5 single-chain Fv (scFv). **Methods.** Two different anti-C5 scFv, one that inhibits both release of C5a and assembly of the TCC (TS-A 12/22) and another that selectively blocks formation of the TCC (TS-A 8), were injected at the onset of AIA. The effects of these scFv on disease severity were evaluated for up to 21 days and compared with the effects of injection of an unrelated scFv. AIA was also established in C6-deficient and C6-sufficient PVG rats to obtain further information on the role of the TCC in this model. **Results.** TS-A 12/22 and TS-A 8 proved to be equally effective in reducing joint swelling, cell counts and tumor necrosis factor alpha levels in synovial lavage fluids, and the degree of histomorphologic changes compared with the effects of the unrelated scFv. TS-A 12/22 and TS-A 8 prevented the deposition of C9 but not that of C3, confirming the ability of the 2 scFv to neutralize C5. Administration of the 2 anti-C5 scFv after AIA onset also reduced disease severity. In C6-deficient rats with AIA, disease activity was reduced markedly compared with that in C6-sufficient rats. **Conclusion.** These 2 human anti-C5 scFv could represent potential therapeutic reagents to be used in patients with rheumatoid arthritis. In addition, the finding that TS-A 8 was as effective as TS-A 12/22 in reducing disease severity suggests that the TCC is mainly responsible for the joint inflammation and damage observed in AIA. (c) 2007, American College of Rheumatology.

? s (antibod? or immunoglobulin? or hybridoma?) (20n) (c5a or ts(w)a12?)

2724863 ANTIBOD?

1064219 IMMUNOGLOBULIN?

61953 HYBRIDOMA?

14523 C5A

50236 TS
 5903 A12?
 4 TS(W)A12?
 S5 1785 (ANTIBOD? OR IMMUNOGLOBULIN? OR HYBRIDOMA?) (20N) (C5A OR
 TS(W)A12?)
 ? s (antibod? or immunoglobulin? or hybridoma?) (20n) (c5a) same (ts(w)a12?)
 2724863 ANTIBOD?
 1064219 IMMUNOGLOBULIN?
 61953 HYBRIDOMA?
 0 C5A) SAME (TS
 0 A12?)
 S6 0 (ANTIBOD? OR IMMUNOGLOBULIN? OR
 HYBRIDOMA?) (20N) (C5A) SAME (TS(W)A12?)
 ? s (antibod? or immunoglobulin? or hybridoma?) (20n) (c5a) and (ts(w)a12?)
 2724863 ANTIBOD?
 1064219 IMMUNOGLOBULIN?
 61953 HYBRIDOMA?
 14523 C5A
 1785 ((ANTIBOD? OR IMMUNOGLOBULIN?) OR HYBRIDOMA?) (20N) C5A
 50236 TS
 5903 A12?
 4 TS(W)A12?
 S7 4 (ANTIBOD? OR IMMUNOGLOBULIN? OR HYBRIDOMA?) (20N) (C5A) AND
 (TS(W)A12?)
 ? rd s7
 S8 2 RD S7 (unique items)
 ? t s8/3/all

8/3/1 (Item 1 from file: 5)
 DIALOG(R)File 5:Biosis Previews(R)
 (c) 2010 The Thomson Corporation. All rts. reserv.

16983076 BIOSIS NO.: 200200576587
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 (Reprint)
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 JOURNAL: European Journal of Immunology 32 (10): p2773-2782 October, 2002
 2002
 MEDIUM: print
 ISSN: 0014-2980
 DOCUMENT TYPE: Article
 RECORD TYPE: Abstract
 LANGUAGE: English

8/3/2 (Item 1 from file: 73)
 DIALOG(R)File 73:EMBASE
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0082118885 EMBASE/Medline No: 2007532089
 Rheumatoid arthritis and the complement system
 Okroj M.; Heinegard D.; Holmdahl R.; Blom A.M.
 Lund University, Department of Laboratory Medicine, University Hospital
 Malmo, Malmo, Sweden
 AUTHOR EMAIL: Anna.Blom@med.lu.se
 CORRESP. AUTHOR/AFFIL: Blom A.M.: Lund University, Department of
 Laboratory Medicine, University Hospital Malmo, Entrance 46, S-205 02 Malmo

, Sweden
CORRESP. AUTHOR EMAIL: Anna.Blom@med.lu.se

Annals of Medicine (Ann. Med.) (Norway) November 15, 2007, 39/7
(517-530)

CODEN: ANMDE ISSN: 0785-3890 eISSN: 1651-2219

PUBLISHER ITEM IDENTIFIER: 780337113

DOI: 10.1080/07853890701477546

DOCUMENT TYPE: Journal; Review RECORD TYPE: Abstract

LANGUAGE: English SUMMARY LANGUAGE: English

NUMBER OF REFERENCES: 139

? s (antibod? or immunoglobulin? or hybridoma?) (20n) (c5a) (20n) (cleavage(w)site)

2724863 ANTIBOD?

1064219 IMMUNOGLOBULIN?

61953 HYBRIDOMA?

14523 C5A

390866 CLEAVAGE

1961132 SITE

S9 3 (ANTIBOD? OR IMMUNOGLOBULIN? OR
HYBRIDOMA?) (20N) (C5A) (20N) (CLEAVAGE(W)SITE)

? rd s9

S10 1 RD S9 (unique items)

? t s10/3/all

10/3/1 (Item 1 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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AUTHOR: Marzari Roberto; Sblattero Daniele; Macor Paolo; Fischetti Fabio;

Gennaro Renato; Marks James D; Bradbury Andrew; Tedesco Francesco

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AUTHOR ADDRESS: Dipartimento di Fisiologia e Patologia, Universita di

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2002

MEDIUM: print

ISSN: 0014-2980

DOCUMENT TYPE: Article

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LANGUAGE: English

?